



Form PTO-1449 U.S. Department of Commerce  
(REV. 2-82) Patent and Trademark Office  
**INFORMATION DISCLOSURE STATEMENT  
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Atty. Docket No.  
32759  
074288.0101

Serial No.  
09/478,737

Applicants  
Wilson et. al.

Examiner  
Murphy

Filing Date  
January 6, 2000

Group Art Unit  
1646

### U.S. PATENT DOCUMENTS

*Exam. Init.	No.	Document No.	Date	Name	Class	Subclass	Filing Date if Appropriate

### FOREIGN PATENT DOCUMENT

		Document No.			Class	SubClass	Translator Yes No

### OTHER DOCUMENTS (including Author, Title Date, Pertinent Pages, Etc.)

A	Peral et al., 1995, "Splicing mutations of the polycystic kidney disease 1 (PKD1) gene induced by intronic deletion," Human Molec. Genet. 4:569-574.
B	The European Polycystic Kidney Disease Consortium, 1994, "The polycystic kidney disease 1 gene encodes a 14kb transcript and lies within a duplicated region on chromosome 16," Cell 77:881-894.
C	Watnick et al., 1997, "An unusual pattern of mutation in the duplicated portion of <i>PKD1</i> is revealed by use of a novel strategy for mutation detection," Human Molecular Genetics 6:1473-1481.
D	Longa et al., 1997, "A large <i>TSC2</i> and <i>PKD1</i> gene deletion is associated with renal and extrarenal signs of autosomal dominant polycystic kidney disease," Nephrology Dialysis Transplantation 12:1900-1907
E	Thomas et al., 1999, "Identification of mutations in the repeated part of the autosomal dominant polycystic kidney disease type 1 gene, PKD1, by long-range PCR," Am. J. Human Genetics 65:39-49.